

1000-875-2000

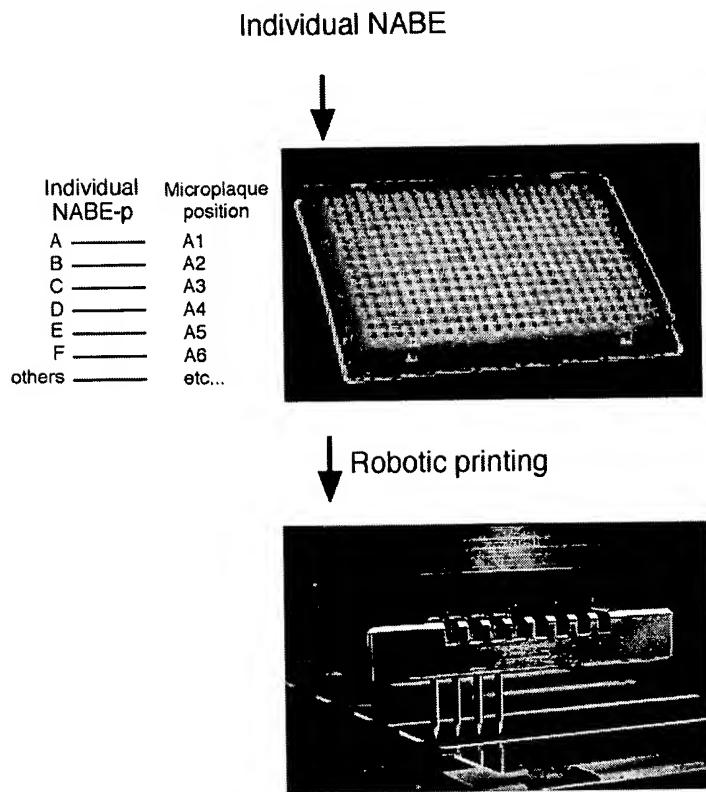


Figure 1. Arraying of individual NABE-p for the microarray fabrication.

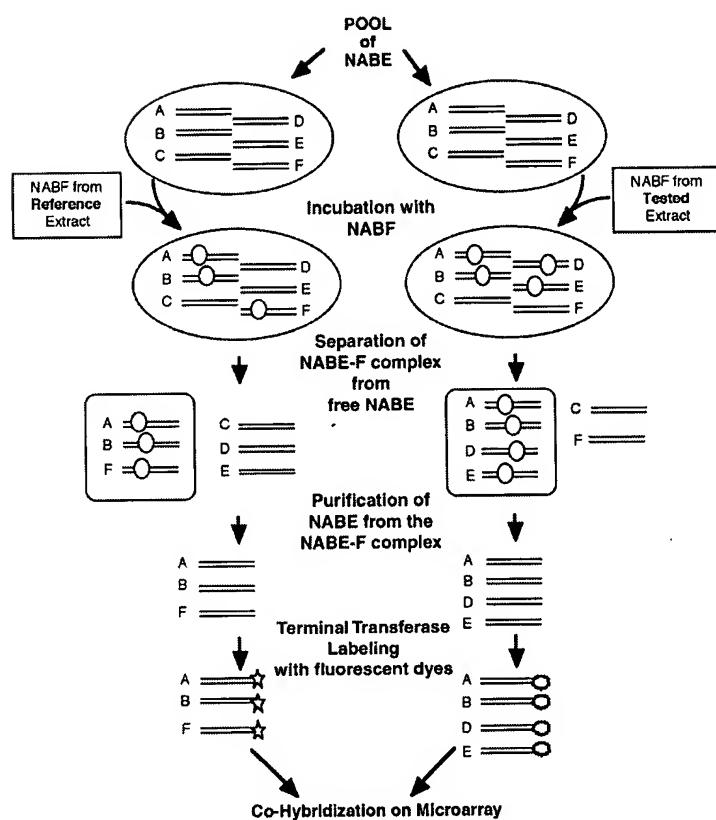
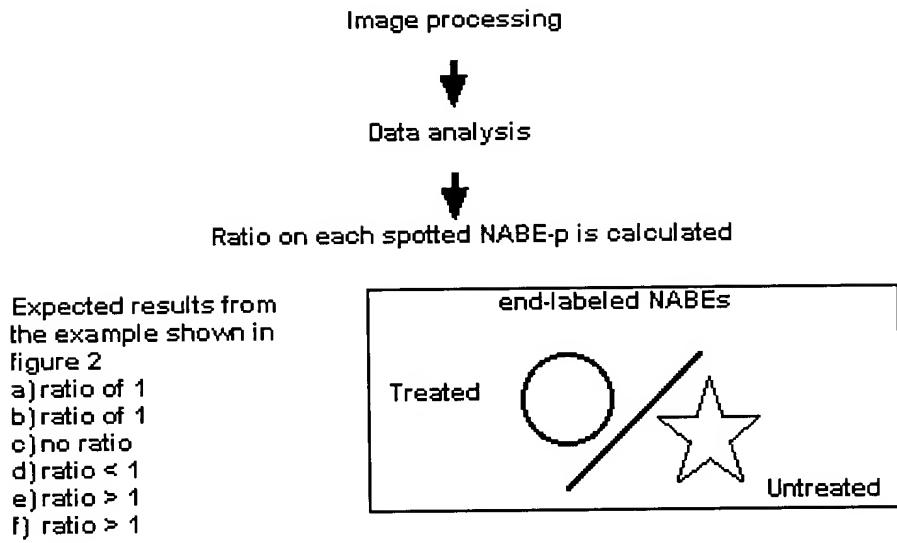
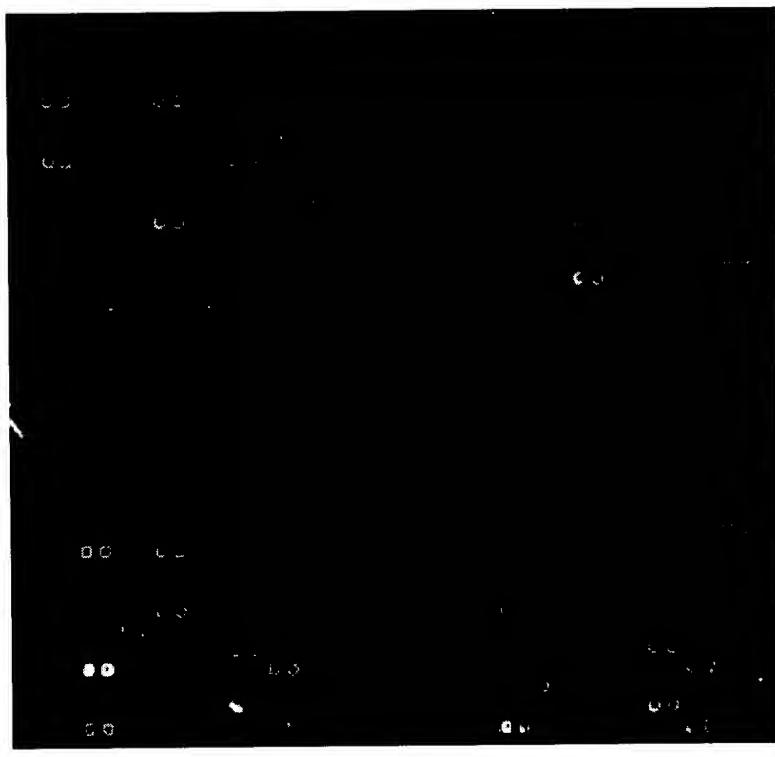


Figure 2. Schematic representation of the analysis of differentially-bound NABEs by DPA.

15  
14  
13  
12  
11  
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9  
8  
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2  
1

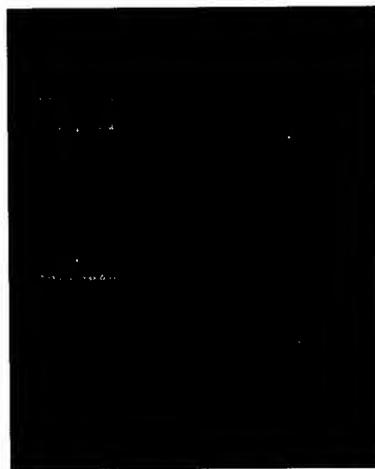


**Figure 3.** Identification of differentially bound NABEs. A ratio of 1 reflects an equivalent DNA-binding activity for a given factor in both conditions tested. A ratio <1 signals a reduced DNA-binding activity while a ratio >1 reflects an increase in such activity. No ratio (no signal) means no factor bound in either condition.

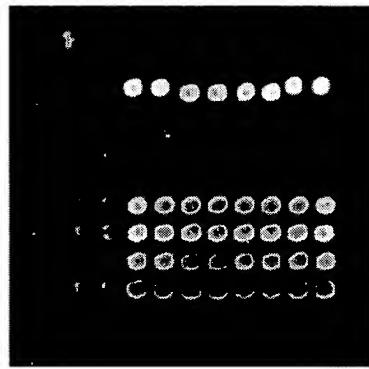


**Figure 4.** A hybridized NABE-p array with Cy5 end-labeled NABEs. 300, 90 and 9 pg of each NABE-p are spotted in duplicate. Non-specific oligos are also spotted as negative controls.

Inventor: LAROSE et al.  
Docket No.: 9555.123USU1  
Title: METHOD FOR SCREENING AND/OR IDENTIFYING PROTEINS THAT BIND TO  
NUCLEIC ACIDS  
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Sheet 5 of 6



**Figure 5.** Hybridization of NABEs retrieved from a shifted NABE-NABF complex to a NABE-p microarray.



**Figure 6.** Co-hybridization of NABEs incubated in K562 cells nuclear extracts and recovered by EMSA to a NABE-p microarray. NABEs incubated with an extract from untreated cells were labeled with Cy3 (green) while NABEs incubated with an extract from TPA-treated cells were labeled with Cy5 (red). Yellow spots reflect equivalent quantities of both types of NABEs.